## What is claimed is:

- 1. A process for preparing a pharmaceutical composition for treating a condition of the central nervous system in a mammalian subject, comprising: reacting gabapentin with tannic acid to produce a pharmaceutically effective amount of gabapentin tannate in solid dosage form wherein the tannic acid component is of either natural or synthetic origin.
- 2. The process of claim 1 including selecting either natural or synthetic tannic acid.
- 3. The process of claim 1 including providing one or more pharmaceutically acceptable excipients.
- 4. A process for preparing a pharmaceutical composition for treating a condition of the central nervous system in a mammalian subject, comprising:

mixing an anti-clumping agent and tannic acid together to form a reaction mixture;

adding gabapentin to said reaction mixture; and adding one or more solvents to said reaction mixture.

5. The process of claim 4, including selecting said solvent from a group consisting of water, purified water, isopropyl alcohol, ethanol, glycerin, propylene glycol, mineral oil and mixtures thereof

6. A process for preparing a pharmaceutical composition for treating a condition of the central nervous system in a mammalian subject, comprising:

mixing one or more anti-clumping agents, tannic acid and gabapentin together either in the presence of one or more solvents or at a suitable temperature so as to produce a pharmaceutically effective amount of gabapentin tannate.

- 7. The process of claim 6, including selecting said solvents from a group consisting of water, purified water, ethanol, isopropyl alcohol, glycerin, propylene glycol, mineral oil and mixtures thereof.
- 8. The process of claim 6, including providing said tannic acid at a weight  $W_1$  and gabapentin at a weight  $W_2$  wherein  $W_1$  is from about 0.05 to about 20 times  $W_2$ .
- 9. The process of claim 8, including selecting said one or more anticlumping agents from a group consisting of magnesium aluminum silicate, xanthan gum, polyvinylpyrrolidone, cellulose compounds, magnesium stearate, colloidal silica, talc, stearic acid, calcium stearate, lactose, mannitol, sucrose and mixtures thereof.
- 10. The process of claim 9, including providing said one or more anticlumping agents at a concentration of from about 0.01 to about 95% by weight of said composition.

- 11. A pharmaceutical composition for treating a condition of the central nervous system in a mammalian subject, comprising as an active ingredient a pharmaceutically effective amount of gabapentin tannate in solid dosage form wherein the tannic acid component is of either natural or synthetic origin.
- 12. The composition of claim 11 further including one or more pharmaceutical excipients.
- 13. The composition of claim 12, wherein said excipients are selected from a group consisting of an anti-clumping agent, a filler, a diluent, a colorant, a sweetening agent, a lubricant, a binding agent, a disintegrating agent, a flavoring agent and mixtures thereof.
- 14. The composition of claim 12, wherein said composition further includes one or more solvents selected from a group consisting of water, purified water, ethanol, isopropyl alcohol, glycerin, propylene glycol, mineral oil and mixtures thereof.
- 15. The composition of claim 12, wherein said one or more excipients are sweetening agents selected from a group consisting of sucrose, saccharin sodium, aspartame, sucralose and mixtures thereof.
- 16. The composition of claim 12, wherein said one or more excipients are anti-clumping agents selected from a group consisting of magnesium aluminum silicate, xanthan gum, polyvinylpyrrolidone, cellulose compounds, magnesium stearate, colloidal silica, talc, stearic acid, calcium stearate, lactose, mannitol,

sucrose and mixtures thereof.

- 17. A method of treating a condition of the central nervous system in a mammalian subject, comprising administering a pharmaceutically effective amount of gabapentin tannate in solid dosage form.
- 18. The method of claim 17 wherein said administering step is performed orally.
- 19. The method of claim 17, including providing between about 0.1 to about 3600 mg of gabapentin in gabapentin tannate salt form.